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AMENDMENTS TO THE CLAIMS II.

This listing of the claims will replace all prior versions, and listings, of claims in the application.

(Currently Amended) A compound of Formula (I) the formula: 1.

$$L_1-X-L_2$$

(I)

wherein:

L_t is a group of formula (a):

$$\begin{array}{c|cccc}
 & R^1 & R^x & B \\
\hline
 & R^2 & R^2 & B
\end{array}$$
(a)

$$\begin{array}{c|c}
R^1 & R^{\times} \\
R^{\times} & B^{\times} - C \\
R^{2} & R^{\times}
\end{array}$$

wherein:

A is phenyl an aryl or a heteroaryl ring;

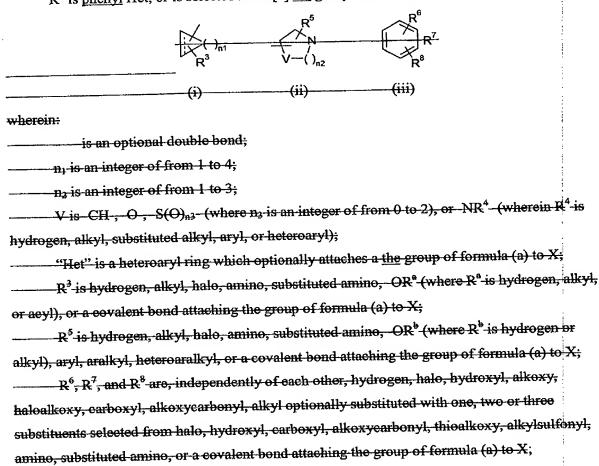
B" is -O-;

Rx is alkyl, alkenyl, or alkynyl, each optionally substituted with 1 to 5 alkoxy or fluoro substituents alkyl, substituted alkyl, alkenyl, substituted alkynyl, substituted alkynyl, acyl, acylamino, aminoacyloxy, aryl, carboxyalkyl, cycloalkyl, substituted cycloalkyl, eycloalkenyl, substituted cycloalkenyl, heteroaryl, heteroaralkyl, alkylsulfonyl, or alkylsulfinyl;

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R1 is hydrogen or alkyl;

R² is phenyl Het, or is selected from [a] the group consisting of formula (i), (ii), and (iii):



K is a bond or an alkylene group;

K" is a bond, C(O), $S(O)_{n4}$ (where n_4 is an integer of from 0 to 2), or an alkylene group optionally substituted with a hydroxyl group; and

B is a piperidine ring heteroeyeloamino or heteroarylamino, which optionally attaches a the group of formula (a) to X;

provided that at least one of the R³, R⁵, R⁶, R⁷, R⁸, "Het", heterocycloamino, or heteroarylamino groups attaches the group of formula (a) to X;

 R^{46} is alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl, or heterocycle; R^{47} is alkyl, substituted alkyl, aryl, acyl, heterocycle, or -COOR⁵⁰ where R^{50} is alkyl; or R^{46} and R^{47} together with the nitrogen atom to which they are attached form a

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heterocycle, which heterocycle is optionally substituted with one or more alkyl, substituted alkyl, alkenyl, substituted alkynyl, alkoxy, substituted alkoxy, cycloalkyl, substituted cycloalkenyl, acyl, acylamino, acyloxy, amino, substituted amino, aminoacyl, substituted cycloalkenyl, acyl, acylamino, acyloxy, amino, substituted amino, aminoacyl, aminoacyloxy, oxyaminoacyl, azido, cyano, halo, hydroxyl, keto, thioketo, carboxyl, carboxyalkyl, thioaryloxy, thioheteroaryloxy, thioheterocyclooxy, thiol, thioalkoxy, substituted thioalkoxy, aryl, aryloxy, heteroaryl, heteroaryloxy, heterocycle, heterocyclooxy, hydroxyamino, alkoxyamino, nitro, -SO-alkyl, -SO-substituted alkyl, -SO-aryl, -SO-heteroaryl, -SO₂-alkyl, -SO₂-substituted alkyl, -SO₂-aryl or -SO₂-heteroaryl;

X is alkylene optionally substituted with one, two, or three hydroxy groups, alkylene wherein one, two, or three carbon atoms have been replaced by an oxygen atom, or an -alkylene-phenylene-alkylene- wherein the phenylene ring is optionally substituted with one or two chloro or fluoro groups:

 L_2 is an organic group comprising at least one primary, secondary or tertiary amine; and X has the formula:

$$X^{a}-Z-(Y^{a}-Z)_{m}-Y^{b}-Z-X^{a}-$$

wherein

m is an integer of from 0 to 20;

X^a at each separate occurrence is selected from the group consisting of O, S, NR,

C(O), C(O)O, C(O)NR, C(S), C(S)O, C(S)NR- or a covalent bond;

Z at each separate occurrence is selected from the group consisting of alkylene, substituted alkylene, substituted eyeloalkylene, alkonylene, substituted alkynylene, substituted alkynylene, substituted eyeloalkenylene, substituted eyeloalkenylene, substituted eyeloalkenylene, substituted eyeloalkenylene, arylene, heteroarylene, heteroarylene, and a covalent bond; and

Y^a and Y^b at each separate occurrence are selected from the group consisting of O,

C(O), OC(O), C(O)O, NR, S(O)n, C(O)NR', NR'C(O), NR'C(O)NR',

NR'C(S)NR', C(=NR') NR', NR'C(=NR'), OC(O) NR', NR'C(O)O, P(O)(OR')O,

OP(O)(OR'), S(O)nCR'R'', S(O)n NR', NR'S(O)n, S.S, and a covalent bond; where n is 0, 1 or 2; and R, R' and R'' at each separate occurrence are selected from the group consisting of hydrogen, alkyl, substituted alkyl, cycloalkyl, substituted eyeloalkyl, alkenyl, substituted alkenyl, substituted eyeloalkenyl, alkenyl, aryl, heteroaryl and alkenyl, cycloalkenyl, substituted alkynyl, aryl, heteroaryl and

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heterocycle; provided at least one of Xo, Yo, Yo, or Z is not a covalent bond;

- or a pharmaceutically acceptable salt; or prodrug thereof. 2. Canceled. Canceled. 3. (Currently Amended) The compound of claim 1 or 2 wherein R1 is hydrogen, methyl, or 4. ethyl. Canceled. 5. Canceled. 6. 7. Canceled. 8. Canceled. Canceled. 9. (Currently Amended) The compound of claim 9 1 wherein Rx is (C1-C6)alkyl, (C2-10. C₆)alkenyl, (C₂-C₆)alkynyl, each optionally substituted with 1 to 3 methoxy, ethoxy or fluoro substituents. (Currently Amended) The compound of claim 9 1 wherein Rx is (C1-C6)alkyl optionally 11. substituted with 1 to 3 methoxy, ethoxy, or fluoro substituents. (Original) The compound of claim 1 wherein Rx is methyl, ethyl, propyl, isopropyl, 12.

 - butyl, isobutyl or secbutyl, optionally substituted with methoxy or ethoxy or with 1 to 3 dr fluoro substituents.

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- 13. (Currently Amended) The compound of claim 1 wherein R^x is methyl, ethyl, methoxymethyl, ethoxymethyl, ethoxyethyl, fluoromethyl, difluoromethyl, trifluoromethyl, or trifluoromethyl, formyl, or acetyl.
- 14. (Currently Amended) The compound of claim 1 or 2 wherein R^x is methyl, ethyl, methoxymethyl, fluoromethyl, difluoromethyl, or trifluoromethyl.
- 15. Canceled.
- 16. Canceled.
- 17. Canceled.
- 18. (Currently Amended) The compound of claim 16 1 wherein B is piperidin-3-yl or piperidin-4-yl wherein the nitrogen at the-1 position optionally attaches the group of formula (a) to X.
- 19. (Original) The compound of claim 1 wherein B taken together with R^x is 4-methylpiperidine-1,4-diyl.
- 20. Canceled.
- 21. (Currently Amended) The compound of claim 20 1 wherein: R^{46} is alkyl or substituted alkyl; R^{47} is alkyl, substituted alkyl, or heterocycle; or R^{46} and R^{47} together with the nitrogen atom to which they are attached form a heterocycle.
- 22. Canceled.
- 23. Canceled.

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- 24. Canceled.
- 25. Canceled.
- 26. Canceled.
- 27. Canceled.
- 28. (Currently Amended) The compound of claim 1 or 2 wherein X is an alkylene group having from 3 to 20 carbon atoms; wherein one or more carbon atoms in the alkylene group is optionally replaced with -O-; and wherein the chain is optionally substituted on carbon with one or more hydroxyl groups.
- 29. (Currently Amended) The compound of claim 1 or 2 wherein X is nonane-1,9-diyl, octane-1,8-diyl, propane-1,3-diyl, 2-hydroxypropane-1,3-diyl, or 5-oxa-nonane-1,9-diyl.
- 30. (Currently Amended) The compound of claim 1 or 2 wherein X has the following formula:

wherein the phenyl ring is optionally substituted with 1, 2, or 3 fluoro groups.

31. (Currently Amended) The compound of claim 1 er 2 wherein X has one of the following formulas:

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- 32. Canceled.
- 33. Canceled.
- 34. (Currently Amended) The compound of claim 1 wherein the compound has the formula having Formula (III):

or a pharmaceutically acceptable salt or prodrug thereof.

- 35. Canceled.
- 36. (Previously Presented) The compound of claim 34 wherein X is an alkylene group having from 3 to 20 carbon atoms; wherein one or more carbon atoms in the alkylene group is optionally replaced with -O-; and wherein the chain is optionally substituted on carbon with one or more hydroxyl groups.
- 37. (Original) The compound of claim 34 wherein X is nonane-1,9-diyl, octane-1,8-diyl,

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propane-1,3-diyl, 2-hydroxypropane-1,3-diyl, or 5-oxa-nonane-1,9-diyl.

38. (Original) The compound of claim 34 wherein X has the following formula:

wherein the phenyl ring is optionally substituted with 1, 2, or 3 fluoro groups.

39. (Previously Presented) The compound of claim 34 wherein X has one of the following formulas:

- 40. Canceled.
- 41. (Currently amended) The compound of claim 20 1 wherein L₂ is a group of formula (d) wherein R⁴⁶ is alkyl that is optionally substituted with from 1 to 5 substituents independently selected from the group consisting of alkoxy, substituted alkoxy, cycloalkyl, substituted cycloalkenyl, acyl, acylamino, acyloxy, amino, aminoacyl, aminoacyloxy, oxyaminoacyl, cyano, halo, hydroxyl, keto, thioketo, carboxyalkyl, thioaryloxy, thioheterocyclooxy, thiol, thioalkoxy, substituted thioalkoxy, heterocycle, heterocyclooxy, hydroxyamino, alkoxyamino, and NR^aR^b, wherein R^a and R^b may be the same or different [and] and are chosen from hydrogen, alkyl, substituted alkyl, cycloalkyl, alkenyl,

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cycloalkenyl, alkynyl, and heterocycle.

- 42. (Currently amended) The compound of claim 20 1 wherein L2 is a group of formula (d) wherein R⁴⁶ is 3-piperidinyl, 4-piperidinyl, or 3-pyrrolidinyl, which R⁴⁶ is optionally substituted with 1 to 3 substituents independently selected from the group consisting of alkoxy, substituted alkoxy, cycloalkyl, substituted cycloalkyl, cycloalkenyl, substituted cycloalkenyl, acyl, acylamino, acyloxy, amino, substituted amino, aminoacyl, aminoacyloxy, oxyaminoacyl, cyano, halo, hydroxyl, keto, thioketo, carboxyalkyl, thioaryloxy, thioheteroaryloxy, thioheterocycloxy, thiol, thioalkoxy, substituted thioalkoxy, heterocycle, heterocyclooxy, hydroxyamino, alkoxyamino, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, and substituted alkynyl.
- 43. (Currently amended) The compound of claim 20 1 wherein R⁴⁶ and R⁴⁷ together with the nitrogen atom to which they are attached form a piperidine or pyrrolidine ring which ring is optionally substituted with 1 to 3 substituents independently selected from the group consisting of alkoxy, substituted alkoxy, cycloalkyl, substituted cycloalkyl, cycloalkenyl, substituted cycloalkenyl, acyl, acylamino, acyloxy, amino, substituted amino, aminoacyl, aminoacyloxy, oxyaminoacyl, cyano, halo, hydroxyl, keto, thioketo, carboxyalkyl, thioaryloxy, thioheterocyclooxy, thioheterocyclooxy, thiol, thioalkoxy, substituted thioalkoxy, heterocycle, heterocyclooxy, hydroxyamino, alkoxyamino, alkyl, substituted alkyl, alkenyl, substituted alkynyl, and substituted alkynyl.
- 44. (Currently amended) The compound of claim 20 1 wherein R⁴⁶ and R⁴⁷ together with the nitrogen atom to which they are attached form a heterocycle that is an aza-crown ether.
- 45. (Original) The compound of claim 44 wherein the aza-crown ether is 1-aza-12-crown-4, 1-aza-15-crown-5, or 1-aza-18-crown-6.
- 46. Canceled.

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- 47. (Currently Amended) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a therapeutically effective amount of a compound of claim 1 or 2.
- 48. Canceled.
- 49. Canceled.
- 50. Canceled.
- 51. (Currently Amended) A The compound of claim 50 having formula (V):

(V)

or a salt thereof.

52. Canceled.

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